

AMENDMENTS TO THE CLAIMS

1. (previously presented) A method for assessing the quantity of a viable microorganism of interest per known quantity of a food product that is present and has been previously applied to the food product in the course of microbially treating the food product, said method comprising:

obtaining a liquid suspension sample comprising different microorganisms removed from a microbial-treated food product and which includes a substantial entirety of a previously applied and viable microorganism of interest from a known quantity of the microbial-treated food product and in which the different microorganisms are suspended in a liquid recovery media of known quantity;

preparing in a manner corresponding to a most probable number model, a series of progressively dilute test samples by combining portions of the liquid suspension sample with a dilution liquid;

incubating the series of progressively dilute test samples for a predetermined period of time under conditions conducive to growth of the microorganism of interest;

conducting a PCR analysis on the series of progressively dilute test samples; and

utilizing the most probable number model to determine the concentration of the viable microorganism of interest present on the food product based on results of the PCR analysis.

2. (previously presented) The method as claimed in claim 1, wherein said food product is a sample of animal feed and said method includes taking the sample of animal feed from a feedpile and transporting the sample to a testing lab in such a way that the sample of the animal feed at the testing laboratory is representative of the condition of the animal feed when the animal feed is to be consumed by animals.
3. (previously presented) The method as claimed in claim 1, wherein said food product is a sample of animal feed and said method includes taking the sample of animal feed from a feedpile at a location where the animal feed is to be consumed by animals.
- 4-6 (canceled)
7. (previously presented) The method as claimed in claim 1, wherein said at least one oligonucleotide hybridizes with a nucleic acid sequence that is indicative of a species of the specific kind of microorganism.
8. (canceled)
9. (currently Amended) The method of claim 1, wherein the series of progressively dilute test samples are divided into multiple portions and after incubating each portion, the PCR analysis detects the presence or absence of the specific kind of microorganism of interest in each incubated portion.
10. (previously presented) The method as claimed in claim 9, wherein the series of progressively dilute test samples are divided into the multiple portions by diluting the test samples and dividing the diluted sample into the multiple portions.

11. (previously presented) The method as claimed in claim 9, wherein the series of progressively dilute test samples are divided into multiple portions by mixing the sample with liquid to produce a fluid mixture, and dividing the fluid mixture into the multiple portions.
12. (currently amended) The method as claimed in claim 1, wherein the PCR analysis comprises ~~the~~ using at least one oligonucleotide to detect the presence or absence of the microorganism of interest in respective portions of the incubated test samples and wherein said PCR analysis includes detecting the presence or absence of a product of hybridization of said at least one ~~oligonucleotide~~ oligonucleotide with a nucleic acid sequence that is indicative of the microorganism of interest.
13. (previously presented) The method as claimed in claim 1, wherein the PCR analysis comprises using two oligonucleotide primers that induce a polymerase chain reaction in the presence of nuclear material of the microorganism of interest, and detecting the presence or absence of a product of the polymerase chain reaction of the two oligonucleotide primers in the presence of the nuclear material of the microorganism of interest.
14. (previously presented) The method as claimed in claim 13, wherein one of the oligonucleotide primers hybridizes with a nucleic acid sequence indicative of the genus of the microorganism of interest, and another of the oligonucleotide primers hybridizes with a nucleic acid sequence indicative of the species of the microorganism.

15. (previously presented) The method as claimed in claim 13, wherein the detecting of the presence or absence of a product of the polymerase chain reaction of the two oligonucleotide primers in the presence of the nuclear material of the microorganism of interest includes performing electrophoresis of polymerase chain reaction products to detect a reaction product having a characteristic molecular length indicative of a polymerase chain reaction of the two oligonucleotide primers in the presence of the nuclear material of microorganism of interest.

16-36 (canceled)

37. (previously presented) The method of claim 1 wherein the microorganism of interest is a probiotic organism.

38. (previously presented) The method of claim 37 wherein the microorganism of interest is selected from the group consisting of *Bacillus subtilis*, *Bifidobacterium adolescentis*, *Bifidobacterium animalis*, *Bifidobacterium bifidum*, *Bifidobacterium infantis*, *Bifidobacterium longum*, *Bifidobacterium thermophilum*, *Lactobacillus acidophilus*, *Lactobacillus agilis*, *Lactobacillus alactosus*, *Lactobacillus alimentarius*, *Lactobacillus amylophilus*, *Lactobacillus amylovorus*, *Lactobacillus animalis*, *Lactobacillus batatas*, *Lactobacillus bavaricus*, *Lactobacillus bifermentans*, *Lactobacillus bifidus*, *Lactobacillus brevis*, *Lactobacillus buchnerii*, *Lactobacillus bulgaricus*, *Lactobacillus catenaeforme*, *Lactobacillus casei*, *Lactobacillus cellobiosus*, *Lactobacillus collinoides*, *Lactobacillus confusus*, *Lactobacillus coprophilus*, *Lactobacillus coryniformis*, *Lactobacillus corynoides*, *Lactobacillus crispatus*, *Lactobacillus curvatus*, *Lactobacillus delbrueckii*, *Lactobacillus desidiosus*, *Lactobacillus divergens*, *Lactobacillus enterii*, *Lactobacillus farciminis*, *Lactobacillus fermentum*, *Lactobacillus frigidus*, *Lactobacillus fructivorans*, *Lactobacillus fructosus*, *Lactobacillus gasseri*, *Lactobacillus halotolerans*, *Lactobacillus helveticus*, *Lactobacillus heterohiochii*, *Lactobacillus hilgardii*, *Lactobacillus hordniae*, *Lactobacillus inulinus*, *Lactobacillus jensenii*,

Lactobacillus jugurti, Lactobacillus kandleri, Lactobacillus kefir, Lactobacillus lactis, Lactobacillus leichmannii, Lactobacillus lindneri, Lactobacillus malefermentans, Lactobacillus mali, Lactobacillus maltaromicus, Lactobacillus minor, Lactobacillus minutus, Lactobacillus mobilis, Lactobacillus murinus, Lactobacillus pentosus, Lactobacillus plantarum, Lactobacillus pseudoplanarum, Lactobacillus reuteri, Lactobacillus rhamnosus, Lactobacillus rogosae, Lactobacillus tolerans, Lactobacillus torquens, Lactobacillus ruminis, Lactobacillus sake, Lactobacillus salivarius, Lactobacillus sanfrancisco, Lactobacillus sharpeae, Lactobacillus trichodes, Lactobacillus vaccinoferus, Lactobacillus viridescens, Lactobacillus vitulinus, Lactobacillus xylophilus, Lactobacillus yamanashiensis, Lactobacillus zeae, Pediococcus acidilactici, Pediococcus pentosaceus, Streptococcus cremoris, Streptococcus discitactis, Streptococcus faecium, Streptococcus intermedius, Streptococcus lactis, Streptococcus thermophilus, and Escherichia coli. Another group of lactate utilizing microorganisms include Propionibacterium freudenreichii, Propionibacterium shermanii, Propionibacterium jensenii, Propionibacterium acidipropionici, Propionibacterium thoenii, Propionibacterium, Megasphaera elsdenii, Selenomonas ruminatum, and Peptostreptococcus asaccharolyticus.

39. (previously presented) The method as claimed in claim 38, wherein the specific kind of probiotic microorganism is a species of Lactobacillus.
40. (previously presented) The method as claimed in claim 38, wherein the specific kind of probiotic microorganism is Lactobacillus acidophilus.
41. (previously presented) The method as claimed in claim 38, wherein the specific kind of probiotic microorganism is Lactobacillus LA-51.